

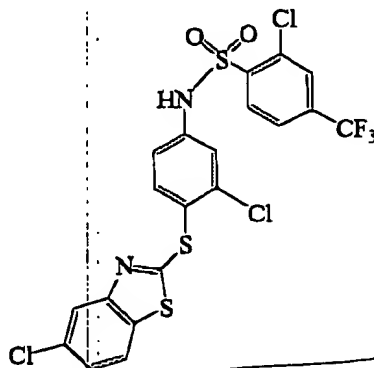
Lawrence R. McGee, *et al.*
Application No.: 09/606,433
Page 2

PATENT

B 1

8 43.

(Amended) A compound of claim 1, having the formula:



B 2

9

46. (Twice Amended) A composition comprising a pharmaceutically acceptable excipient and a compound of any one of claims 1, 2, ~~4-8~~³⁻⁷, and ~~43~~⁸.

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47. (Twice Amended) A method for treating a condition mediated by PPAR γ in a host, said method comprising administering to said host an efficacious amount of a compound of any one of claims 1, 2, ~~4-8~~³⁻⁷, and ~~43~~⁸.

B 3

28 66.

(Twice Amended) A compound of any of claims 1, 2, ~~4-8~~³⁻⁷, and ~~43~~⁸, wherein said compound is a modulator of PPAR γ .

REMARKS

Claims 1, 2, 4-8, 43, 46-53, and 55-67 are presented for examination. Claims 43, 46, 47, and 66 are amended. Claim 44 is canceled without prejudice.

Amendments to the Claims

The amendments to the claims add no new matter.

Claim 43 was amended to delete a recital of one of the two members of the Markush group.

Claims 46, 47 and 66 were amended to remove the recital of canceled claim 44 from their multidependency recitals.

Lawrence R. McGee, *et al.*
Application No.: 09/606,433
Page 3

PATENT

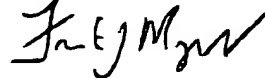
As the Amendments to the claims add no new matter, Applicants respectfully request their entry.

CONCLUSION

In view of the foregoing, Applicants believe all claims now pending in this Application are in condition for allowance. The issuance of a formal Notice of Allowance at an early date is respectfully requested.

If the Examiner believes a telephone conference would expedite prosecution of this application, please telephone the undersigned at 925-472-5012.

Respectfully submitted,



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WC 9046833 v1

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Lawrence R. McGee, *et al.*
Application No.: 09/606,433
Page 4

PATENT

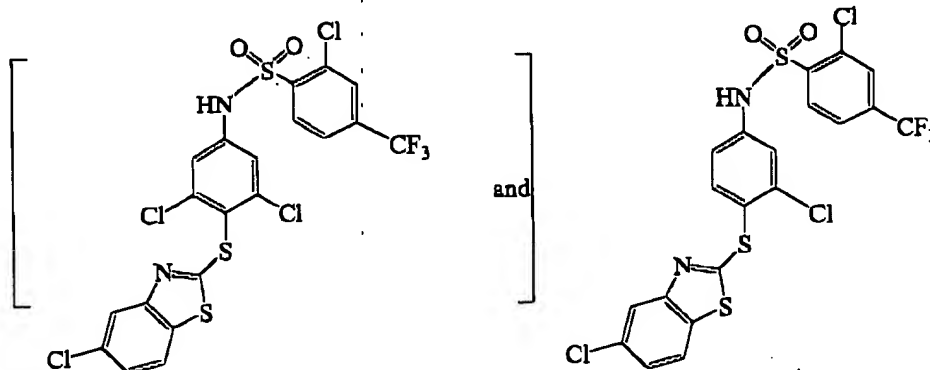
VERSION WITH MARKINGS TO SHOW CHANGES MADE

In the claims:

Claim 44 has been canceled without prejudice.

Claims 43, 46, 47, and 66 have been amended as follows:

43. (Amended) A compound of claim 1, [selected from the group consisting of] having the formula:



46. (Twice Amended) A composition comprising a pharmaceutically acceptable excipient and a compound of any one of claims 1, 2, 4-8, and 43[, and 44].

47. (Twice Amended) A method for treating a condition mediated by PPAR γ in a host, said method comprising administering to said host an efficacious amount of a compound of any one of claims 1, 2, 4-8, and 43[, and 44].

66. (Twice Amended) A compound of any of claims 1, 2, 4-8, and 43[, and 44], wherein said compound is a modulator of PPAR γ .

WC 9046833 v1